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November 13, 1997

Assistant Commissioner for Patents
Washington, DC 20231
Box Patent Application

TRANSMITTAL LETTER SMALL ENTITY APPLICATION

Dear Sir:

Please find enclosed a patent application as follows:

Applicant(s): Moungi Bawendi, Klavs F. Jensen, Bashire O. Dabbousi, Xavier Rodriguez-Viejo and Frederic Victor Mikulec

Title: HIGHLY LUMINESCENT COLOR-SELECTIVE MATERIALS

No. Pages Specification: 16 No. Pages Claims: 4; No. Sheets Dwgs.: 5; No. Pages Abstract: 1;
Unexecuted Formal Papers: Small Entity Statement; Combined Declaration and Power of Attorney,
Assignment;

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Additional Fees:

Total Number of Claims in excess of 20 times \$11: 28-20=8 88.00

Number of independent claims minus 3 times \$41: 3-3=0 0.00

Multiple Dependent Claims (\$135): 135.00

Total Filing Fee \$618.00

Enclosed please find a check in the amount of \$618 to cover the appropriate filing fees. Also enclosed are an Information Disclosure Statement, Form PTO-1449 and cited art. Additionally, please charge any further fees, and credits, to our Deposit Account No. 03-1721.

If this application is found otherwise to be INCOMPLETE, or if at any time it appears that a TELEPHONE CONFERENCE with counsel would helpfully advance prosecution, please telephone the undersigned at any time.

Kindly acknowledge receipt of the foregoing application by returning the self-addressed postcards.

Respectfully submitted,

Mary Rose Scozzafava

Mary Rose Scozzafava, Reg. No. 36,268

JOINT

**APPLICATION
FOR
UNITED STATES LETTER PATENT**

TO THE COMMISSIONER OF PATENTS AND TRADEMARKS:

BE IT KNOWN, that We, Moungi Bawendi, Klavs F. Jensen, Bashir O. Dabbousi, Xavier Rodriguez-Viejo and Frederic Victor Mikulec have invented certain new useful improvements in HIGHLY LUMINESCENT COLOR-SELECTIVE MATERIALS of which the following is a specification:

HIGHLY LUMINESCENT COLOR-SELECTIVE MATERIALS

DS1 349876 2

Field of the Invention

5 This invention relates to luminescent nanocrystalline materials which emit visible light over a very narrow range of wavelengths. The invention further relates to materials which emit visible light over a narrow range tunable over the entire visible spectrum.

Background of the Invention

10 Semiconductor nanocrystallites (quantum dots) whose radii are smaller than the bulk exciton Bohr radius constitute a class of materials intermediate between molecular and bulk forms of matter. Quantum confinement of both the electron and hole in all three dimensions leads to an increase in the effective band gap of the material with decreasing crystallite size. Consequently, both the optical absorption and emission of quantum dots
15 shift to the blue (higher energies) as the size of the dots gets smaller.

 Bawendi and co-workers have described a method of preparing monodisperse semiconductor nanocrystallites by pyrolysis of organometallic reagents injected into a hot coordinating solvent (*J. Am. Chem. Soc.*, **115**:8706 (1993)). This permits temporally discrete nucleation and results in the controlled growth of macroscopic quantities of
20 nanocrystallites. Size selective precipitation of the crystallites from the growth solution provides crystallites with narrow size distributions. The narrow size distribution of the quantum dots allows the possibility of light emission in very narrow spectral widths.

 Although semiconductor nanocrystallites prepared as described by Bawendi and co-workers exhibit near monodispersity, and hence, high color selectivity, the
25 luminescence properties of the crystallites are poor. Such crystallites exhibit low photoluminescent yield, that is, the light emitted upon irradiation is of low intensity. This is due to energy levels at the surface of the crystallite which lie within the energetically forbidden gap of the bulk interior. These surface energy states act as traps for electrons and holes which degrade the luminescence properties of the material.

In an effort to improve photoluminescent yield of the quantum dots, the nanocrystallite surface has been passivated by reaction of the surface atoms of the quantum dots with organic passivating ligands, so as to eliminate forbidden energy levels. Such passivation produces an atomically abrupt increase in the chemical potential at the interface of the semiconductor and passivating layer (See, A.P. Alivisatos, *J. Phys. Chem.* **100**:13226 (1996)). Bawendi et al. (*J. Am. Chem. Soc.*, **115**:8706 (1993)) describe CdSe nanocrystallites capped with organic moieties such as tri-*n*-octyl phosphine (TOP) and tri-*n*-octyl phosphine oxide (TOPO) with quantum yields of around 5-10%.

Passivation of quantum dots using inorganic materials also has been reported.

Particles passivated with an inorganic coating are more robust than organically passivated dots and have greater tolerance to processing conditions necessary for their incorporation into devices. Previously reported inorganically passivated quantum dot structures include CdS-capped CdSe and CdSe-capped CdS (Tian et al., *J. Phys. Chem.* **100**:8927 (1996)); ZnS grown on CdS (Youn et al., *J. Phys. Chem.* **92**:6320 (1988)); ZnS on CdSe and the inverse structure (Kortan et al., *J. Am. Chem. Soc.* **112**:1327 (1990)); and SiO₂ on Si (Wilson et al., *Science* **262**:1242 (1993)). These reported quantum dots exhibit very low quantum efficiency and hence are not commercially useful in light emitting applications.

M.A. Hines and P. Guyot-Sionnest report the preparation of ZnS-capped CdSe nanocrystallites which exhibited a significant improvement in luminescence yields of up to 50% quantum yield at room temperature (*J. Phys. Chem.* **100**:468 (1996)). However, the quality of the emitted light remained unacceptable because of the large size distribution (12-15% rms) of the core of the resulting capped nanocrystallites. The large size distribution resulted in light emission over a wide spectral range. In addition, the reported preparation method does not allow control of the particle size obtained from the process and hence does not allow control of color.

Danek *et al.* report the electronic and chemical passivation of CdSe nanocrystals with a ZnSe overlayer (*Chem. Materials* **8**:173 (1996)). Although it might be expected that such ZnSe-capped CdSe nanocrystallites would exhibit as good as or better quantum

yield than the ZnS analogue due to the better unit cell matching of ZnSe, in fact, the resulting material showed only disappointing improvements in quantum efficiency (≤ 0.4 % quantum yield).

Thus there remains a need for semiconductor nanocrystallites capable of light
5 emission with high quantum efficiencies throughout the visible spectrum, which possess a narrow particle size (and hence with narrow photoluminescence spectral range).

It is the object of the invention to provide semiconductor nanocrystallites which overcome the limitations of the prior art and which exhibit high quantum yields with photoluminescence emissions of high spectral purity.

10

Summary of the Invention

In one aspect of the invention, a coated nanocrystal capable of light emission includes a substantially monodisperse core selected from the group consisting of CdX, where X = S, Se, Te; and an overcoating of ZnY, where Y = S, Se, and mixtures thereof
15 uniformly deposited thereon, said coated core characterized in that when irradiated the particles emit light in a narrow spectral range of no greater than about 40 nm at full width half max (FWHM). In some embodiments, the narrow spectral range is selected from the spectrum in the range of about 470 nm to about 620 nm and the particle size of the core is selected from the range of about 20 Å to about 125 Å.

20 In other embodiments of the invention, the coated nanocrystal is characterized in that the nanocrystal exhibits less than a 10% and preferably less than 5%, rms deviation in diameter of the core. The nanocrystal preferably exhibits photoluminescence having quantum yields of greater than 30%, and most preferably in the range of about 30 to 50%.

25 In another embodiment of the invention, the overcoating comprises one to two monolayers of ZnY. The nanocrystal may further comprise an organic layer on the nanocrystal outer surface. The organic layer may be comprised of moieties selected to provide compatibility with a suspension medium, such as a short-chain polymer terminating in a moiety having affinity for a suspending medium, and moieties which

demonstrate an affinity to the quantum dot surface. The affinity for the nanocrystal surface promotes coordination of the organic compound to the quantum dot outer surface and the moiety with affinity for the suspension medium stabilizes the quantum dot suspension.

5 In another aspect of the invention, a method of preparing a coated nanocrystal capable of light emission includes introducing a substantially monodisperse first semiconductor nanocrystal and a precursor capable of thermal conversion into a second semiconductor material into a coordinating solvent. The coordinating solvent is maintained at a temperature sufficient to convert the precursor into the second
10 semiconductor material yet insufficient to substantially alter the monodispersity of the first semiconducting nanocrystal and the second semiconductor material has a band gap greater than the first semiconducting nanocrystal. An overcoating of the second semiconductor material is formed on the first semiconducting nanocrystal.

 In one embodiment of the invention, the monodispersity of the nanocrystal is
15 monitored during conversion of the precursor and overcoating of the first semiconductor nanocrystal. In another embodiment, an organic overcoating is present on the outer nanocrystal surface, obtained by exposing the nanocrystal to an organic compound having affinity for the nanocrystal surface, whereby the organic compound displaces the coordinating solvent.

20 In addition to having higher quantum efficiencies, ZnS overcoated particles are more robust than organically passivated nanocrystallites and are potentially more useful for optoelectronic devices. The (CdSe)ZnS dots of the invention may be incorporated into electroluminescent devices (LEDs). In addition, the (CdSe)ZnS dots of the invention may exhibit cathodoluminescence upon excitation with both high and low voltage
25 electrons and may be potentially useful in the production of alternating current thin film electroluminescent devices (ACTFELD). In the naming convention used herein to refer to capped nanocrystallites, the compound found within parentheses represents the core compound (i.e. the bare "dot"), while the compound which follows represents the overcoated passivation layer.

These and other features and advantages of the invention are set forth in the description of the invention, which follows.

Brief Description of the Drawing

5 The invention is described with reference to the figures, which are presented for the purpose of illustration only, and in which:

Figure 1 shows the absorption spectra of CdSe dots with diameters measuring (a) 23 Å, (b) 42 Å, (c) 48 Å and (d) 55 Å before (dashed lines) and after (solid lines) overcoating with 1-2 monolayers of ZnS

10 Figure 2 shows the room temperature photoluminescence (PL) spectra of the samples of Figure 1 before (dashed lines) and after (solid lines) overcoating with ZnS;

Figure 3 is a color photograph which illustrates the wide spectral range of luminescence from and color purity of the (CdSe)ZnS composite quantum dots of the present invention;

15 Figure 4 shows the progression of the absorption spectra for (CdSe)ZnS quantum dots with ZnS coverages of approximately 0, 0.65, 1.3, 2.6 and 5.3 monolayers; and

Figure 5 shows the evolution of the PL for ~40 Å diameter (CdSe)ZnS dots of Figure 4 with varying ZnS coverage,

Detailed Description of the Invention

20 The present invention is directed to the preparation of a series of room temperature, highly luminescent ZnS-capped CdSe ((CdSe)ZnS) nanocrystallites having a narrow particle size distribution. Nanocrystallites of the present invention exhibit high quantum yields greater than about 30% and preferably in the range of about 30-50% and
25 a narrow band edge luminescence spanning most of the visible spectrum from 470 nm to 625 nm. The core of the nanocrystallites is substantially monodisperse. By monodisperse, as that term is used herein, it is meant a colloidal system in which the suspended particles have substantially identical size and shape. For the purposes of the present invention, monodisperse particles deviate less than 10% in rms diameter in the
30 core, and preferably less than 5% in the core.

When capped quantum dots of the invention are illuminated with a primary light source, a secondary emission of light occurs of a frequency that corresponds to the band gap of the semiconductor material used in the quantum dot. As previously discussed, the band gap is a function of the size of the nanocrystallite. As a result of the narrow size distribution of the capped nanocrystallites of the invention, the illuminated quantum dots emit light of a narrow spectral range resulting in high purity light. Spectral emissions in a narrow range of no greater than about 60 nm, preferably 40 nm and most preferably 30 nm at full width half max (FWHM) are observed.

The present invention also is directed to a method of making capped quantum dots with a narrow particle size distribution. The capped quantum dots of the invention may be produced using a two step synthesis in which a size selected nanocrystallite is first synthesized and then overcoated with a passivation layer of a preselected thickness. In preferred embodiments, processing parameters such as reaction temperature, extent of monodispersity and layer thickness may be monitored during crystal growth and overcoating to provide a coated quantum dot of narrow particle size distribution, high spectral purity and high quantum efficiency. "Quantum yield" as that term is used herein, means the ratio of photons emitted to that absorbed, e.g., the photoluminescence quantum yield.

The method is described for a (CdSe)ZnS quantum dot, but it is understood that the method may be applied in the preparation of a variety of known semiconductor materials. The first step of a two step procedure for the synthesis of (CdSe)ZnS quantum dots involves the preparation of nearly monodisperse CdSe nanocrystallites. The particles range in size from about 23Å to about 55Å with a particle size distribution of about 5-10 % . These dots are referred to as "bare" dots. The CdSe dots are obtained using a high temperature colloidal growth process, followed by size selective precipitation.

The high temperature colloidal growth process is accomplished by rapid injection of the appropriate organometallic precursor into a hot coordinating solvent to produce a temporally discrete homogeneous nucleation. Temporally discrete nucleation is attained by a rapid increase in the reagent concentration upon injection, resulting in an abrupt

supersaturation which is relieved by the formation of nuclei and followed by growth on the initially formed nuclei. Slow growth and annealing in the coordinating solvent results in uniform surface derivatization and regularity in the core structure.

5 Injection of reagents into the hot reaction solvent results in a short burst of homogeneous nucleation. The depletion of reagents through nucleation and the sudden temperature drop associated with the introduction of room temperature reagents prevents further nucleation. The solution then may be gently heated to reestablish the solution temperature. Gentle reheating allows for growth and annealing of the crystallites. The higher surface free energy of the small crystallites makes them less stable with respect to
10 dissolution in the solvent than larger crystallites. The net result of this stability gradient is the slow diffusion of material from small particles to the surface of large particles (“Ostwald ripening”). Growth of this kind results in a highly monodisperse colloidal suspension from systems which may initially be highly polydisperse.

Both the average size and the size distribution of the crystallites in a sample are
15 dependent on the growth temperature. The growth temperature necessary to maintain steady growth increases with increasing average crystal size. As the size distribution sharpens, the temperature may be raised to maintain steady growth. As the size distribution sharpens, the temperature may be raised in 5-10 °C increments to maintain steady growth. Conversely, if the size distribution begins to spread, the temperature may
20 be decreased 5-10 °C to encourage Ostwald ripening and uniform crystal growth. Generally, nanocrystallites 40 Angstroms in diameter can be grown in 2-4 hours in a temperature range of 250-280 °C. Larger samples (60 Angstroms or more) can take days to grow and require temperatures as high as 320 °C. The growth period may be shortened significantly (e.g., to hours) by using a higher temperature or by adding
25 additional precursor materials.

Size distribution during the growth stage of the reaction may be approximated by monitoring the absorption line widths of the particles. Modification of the reaction temperature in response to changes in the absorption spectrum of the particles allows the maintenance of a sharp particle size distribution during growth. It is also contemplated

that reactants could be added to the nucleation solution during crystal growth to grow larger crystals.

The particle size distribution may be further refined by size selective precipitation. In a preferred embodiment, this may be accomplished by manipulation of solvent composition of the nanocrystallite suspension.

The CdSe nanocrystallites are stabilized in solution by the formation of a lyophilic coating of alkyl groups on the crystallite outer surface. The alkyl groups are provided by the coordinating solvent used during the growth period. The interparticle repulsive force introduced by the lyophilic coating prevents aggregation of the particles in solution. The effectiveness of the stabilization is strongly dependent upon the interaction of the alkyl groups with the solvent. Gradual addition of a non-solvent will lead to the size-dependent flocculation of the nanocrystallites. Non-solvents are those solvents in which the groups which may be associated with the crystallite outer surface show no great affinity. In the present example, where the coordinating group is an alkyl group, suitable non-solvents include low molecular weight alcohols such as methanol, propanol and butanol. This phenomenon may be used to further narrow the particle size distribution of the nanocrystallites by a size-selective precipitation process. Upon sequential addition of a non-solvent, the largest particles are the first to flocculate. The removal of a subset of flocculated particles from the initial solution results in the narrowing of the particle size distribution in both the precipitate and the supernatant.

A wealth of potential organometallic precursors and high boiling point coordinating solvents exist which may be used in the preparation of CdSe dots. Organometallic precursors are selected for their stability, ease of preparation and clean decomposition products and low cracking temperatures. A particularly suitable organometallic precursor for use as a Cd source include alkyl cadmium compounds, such as CdMe_2 . Suitable organometallic precursors for use as a Se source include, bis(trimethylsilyl)selenium ($(\text{TMS})_2\text{Se}$), (tri-*n*-octylphosphine)selenide (TOPSe) and trialkyl phosphine selenides, such as (tri-*n*-butylphosphine)selenide (TBPSe). Other suitable precursors may include both cadmium and selenium in the same molecule. Alkyl phosphines and alkyl phosphine oxides may be used as a high boiling coordinating

solvent; however, other coordinating solvents, such as pyridines, furans, and amines may also be suitable for the nanocrystallite production.

The preparation of monodisperse CdSe quantum dots has been described in detail in Murray *et al.* (*J. Am. Chem. Soc.*, **115**:8706 (1993)), which is hereby incorporated in its entirety by reference.

Next, the CdSe particles are overcoated by introducing a solution containing zinc and sulfur precursors in a coordinating solvent (e.g., TOP) into a suspension of CdSe nanocrystallites at the desired temperature. The temperature at which the dots are overcoated is related to the quality of the resultant composite particle. Overcoating the CdSe particles at relatively higher temperatures may cause the CdSe seed crystals to begin to grow via Ostwald ripening and deterioration of the size distribution of the particles leading to broader spectral line widths. Overcoating the particles at relatively low temperatures could lead to incomplete decomposition of the precursors or to reduced crystallinity of the ZnS shell. An ideal growth temperature may be determined for each CdSe core size to ensure that the size distribution of the cores remains constant and that shells with a high degree of crystallinity are formed. In preferred embodiments, CdSe crystallites are overcoated using diethyl zinc and hexamethyldisilathiane as the zinc and sulfur precursors. CdSe crystallites having a diameter in the range of about 23Å-30Å are overcoated at a temperature in the range of about 135-145 °C, and preferably about 140 °C. Similarly, nanocrystallites having a diameter of about 35Å, 40Å, 48Å, and 55Å, respectively, are overcoated at a temperature of about 155-165 °C, and preferably about 160 °C, 175-185 °C and preferably about 180 °C, about 195-205 °C, and preferably about 200 °C, and about 215-225 °C, and preferably about 220 °C, respectively. The actual temperature ranges may vary, dependent upon the relative stability of the precursors and the crystallite core and overlayer composition. These temperature ranges may need to be modified 10-20 °C, depending upon the relative stability of the precursors. For example, when the more stable trialkyl phosphine chalcogenides (like TOPSe) are used, higher temperatures are employed. The resulting (CdSe)ZnS composite particles are also passivated with TOPO/TOP on their outermost surface.

The ZnS precursor solution concentration and the rate of its addition to the CdSe particles is selected to promote heterogeneous growth of ZnS onto the CdSe nuclei instead of homogeneous nucleation to produce ZnS particles. Conditions favoring heterogeneous growth include dropwise addition, e.g., 1-2 drops/second, of the ZnS precursor solution to the CdSe solution and maintenance of the ZnS precursor solution at low concentrations. Low concentrations typically range from 0.0005-0.5 M. In some preferred embodiments, it may be desirable to include a final purification step in which the overcoated dots are subjected to size selective precipitation to further assure that mainly only (CdSe)ZnS composite particles are present in the final product.

In other embodiments, it may be desirable to modify the crystallite outer surface to permit formation of stable suspensions of the capped quantum dots. The outer surface of the nanocrystal includes an organic layer derived from the coordinating solvent used during the capping layer growth process. The crystallite surface may be modified by repeated exposure to an excess of a competing coordinating group. For example, a dispersion of the capped quantum dot may be treated a coordinating organic compound, such as pyridine, to produce crystallites which dispersed readily in pyridine, methanol, and aromatics but no longer dispersed in aliphatics. Such a surface exchange process may be carried out using a variety of compounds which are capable of coordinating or bonding to the outer surface of the capped quantum dot, such as by way of example, phosphines, thiols, amines and phosphates. In other embodiments, the capped quantum dots may be exposed to short chained polymers which exhibit an affinity for the capped surface on one and which terminate in a moiety having an affinity for the suspension or dispersion medium. Such affinity improves the stability of the suspension and discourages flocculation of the capped quantum dots.

The synthesis described above produces overcoated quantum dots with a range of core and shell sizes. Significantly, the method of the invention allows both the size distribution of the nanocrystallites and the thickness of the overcoating to be independently controlled. Figure 1 shows the absorption spectra of CdSe dots with a particle size distribution of (a) 23 Å, (b) 42 Å, (c) 48 Å and (d) 55 Å in diameter before (dashed lines) and after (solid lines) overcoating with 1-2 monolayers of ZnS. By

“monolayer” as that term is used herein, it is meant a shell of ZnS which measures 3.1 Å (the distance between consecutive planes along the [002] axis in the bulk wurtzite ZnS) along the major axis of the prolate shaped dots. The absorption spectra represents the wavelength and intensity of absorption of light which is absorbed by the quantum dot.

5 Figure 1 indicates a small shift in the absorption spectra to the red (lower energies) after overcoating due to the partial leakage of the exciton into the ZnS matrix. This red shift is more pronounced in smaller dots where the leakage of the exciton into the ZnS shell has a more dramatic effect on the confinement energies of the charge carriers.

Figure 2 shows the room temperature photoluminescence spectra (PL) of the
10 samples shown in Figure 1 before (dashed lines) and after (solid lines) overcoating with ZnS. The PL quantum yield increases from 5-15% for bare dots to values ranging from 30% to 50% for dots passivated with ZnS. The PL spectra are much more intense due to their higher quantum yield of (a) 40%, (b) 50%, (c) 35% and (d) 30%, respectively. The quantum yield reaches a maximum value with the addition of approximately 1.3
15 monolayers of ZnS. A decrease in quantum yields at higher ZnS coverages may be due to the formation of defects in the ZnS shell.

Figure 3 is a color photograph which demonstrates the wide spectral range of luminescence from the (CdSe)ZnS composite quantum dots of the present invention. The photograph shows six different samples of ZnS overcoated CdSe dots dispersed in dilute
20 hexane solutions and placed in identical quartz cuvettes. The samples were irradiated with 356 nm ultraviolet light from a uv lamp in order to observe luminescence from all solutions at once. As the size of the CdSe core increased, the color of the luminescence shows a continuous progression from the blue through the green, yellow, orange to red. Their PL peaks occur at (going from right to left in Figure 3) (a) 470 nm, (b) 480 nm, (c)
25 520 nm, (d) 560 nm, (e) 594 nm and (f) 620 nm. In contrast, in the smallest sizes of bare TOPO-capped dots, the color of the PL is normally dominated by broad deep trap emissions and appears as faint white light.

In order to demonstrate the effect of ZnS passivation on the optical and structural properties of CdSe dots, a large quantity of ~40 Å ($\pm 10\%$) diameter CdSe dots were
30 overcoated with varying amounts of Zn and S precursors under identical temperatures

and variable times. The result was a series of samples with similar CdSe cores, but with varying ZnS shell thicknesses. Figure 4 shows the progression of the absorption spectrum for these samples with ZnS coverages of approximately 0 (bare TOPO capped CdSe), 0.65, 1.3, 2.6 and 5.3 monolayers. The right hand side of the figure shows the long wavelength region of the absorption spectra showing the lowest energy optical transitions. The spectra demonstrate an increased red-shift with the thicker ZnS overcoating as well as a broadening of the first peak in the spectra due to increased polydispersity of shell thicknesses. The left hand side of the spectra show the ultra-violet region of the spectra indicating an increased absorption at higher energies with increasing ZnS thickness due to direct absorption into the higher ZnS band gap ZnS shell.

The evolution of the PL for the same ~ 40 Å diameter CdSe dots with ZnS coverage is displayed in Figure 5. As the coverage of ZnS on the CdSe surface increases one observes a dramatic increase in the fluorescence quantum yield followed by a steady decline after ~ 1.3 monolayers of ZnS. The spectra are red shifted (slightly more than the shift in the absorption spectra) and show an increased broadening at higher coverages. The inset to Figure 5 charts the evolution of the quantum yield for these dots as a function of the ZnS shell thickness. For this particular sample, the quantum yield started at 15% for the bare TOPO capped CdSe dots and increased with the addition of ZnS approaching a maximum value of 50% at approximately ~ 1.3 monolayer coverage. At higher coverages, the quantum yield began to decrease steadily until it reached a value of about 30% at about 5 monolayers coverage.

Although the invention has been described with reference to the preparation and performance of CdSe(ZnS), it will be readily apparent that the method of preparation may be used to obtain monodisperse overcoated quantum dots with various combinations of nanocrystallite core and overcoating. The method of the invention permits the preparation of a variety of capped nanocrystals having a very narrow particle size distribution and exhibiting improvements in color purity and intensity of their photoluminescent emissions. It is contemplated that a variety of cadmium chalcogenides, for example, CdX, where X = S, Se, Te may be prepared and overcoated according to the

method of the invention. It is further contemplated that the overcoating may be varied and may include, by way of example only, ZnS, ZnSe, CdS and mixtures thereof.

The invention is described with reference to the following examples, which are presented for the purpose of illustration and which are not intended to be limiting of the invention, the scope of which is set forth in the claims which follow this specification.

Example 1. Preparation of CdSe. Trioctylphosphine oxide (TOPO, 90% pure) and trioctylphosphine (TOP, 95% pure) were obtained from Strem and Fluka, respectively. Dimethyl cadmium (CdMe_2) and diethyl zinc (ZnEt_2) were purchased from Alfa and Fluka, respectively, and both materials were filtered separately through a 0.2 μm filter in an inert atmosphere box. Trioctylphosphine selenide was prepared by dissolving 0.1 mols of Se shot in 100ml of TOP thus producing a 1M solution of TOPSe. Hexamethyl(disilathiane) (TMS_2S) was used as purchased from Aldrich. HPLC grade n-hexane, methanol, pyridine and n-butanol were purchased from EM Sciences.

The typical preparation of TOP/TOPO capped CdSe nanocrystallites follows. TOPO (30g) was placed in a flask and dried under vacuum (~ 1 Torr) at 180 °C for 1 hour. The flask was then filled with nitrogen and heated to 350 °C. In an inert atmosphere drybox the following injection solution was prepared: CdMe_2 (200 microliters, 2.78 mmol), 1 M TOPSe solution (4.0 mL, 4.0 mmol), and TOP (16 mL). The injection solution was thoroughly mixed, loaded into a syringe, and removed from the drybox.

The heat was removed from the reaction flask and the reagent mixture was delivered into the vigorously stirring TOPO with a single continuous injection. This produces a deep yellow/orange solution with a sharp absorption feature at 470-500 nm and a sudden temperature decrease to ~ 240 °C. Heating was restored to the reaction flask and the temperature was gradually raised to 260-280 °C.

Aliquots of the reaction solution were removed at regular intervals (5-10 min) and absorption spectra taken to monitor the growth of the crystallites. The best samples were prepared over a period of a few hours steady growth by modulating the growth temperature in response to changes in the size distribution, as estimated from the sharpness of the features in the absorption spectra. The temperature was lowered 5-10 °C

in response to an increase in the size distribution. Alternatively, the reaction can also be stopped at this point. When growth appears to stop, the temperature is raised 5-10 °C. When the desired absorption characteristics were observed, the reaction flask was allowed to cool to ~60 °C and 20 mL of butanol were added to prevent solidification of the TOPO. Addition of a large excess of methanol causes the particles to flocculate. The flocculate was separated from the supernatant liquid by centrifugation; the resulting powder can be dispersed in a variety of organic solvents (alkanes, ethers, chloroform, tetrahydrofuran, toluene, etc.) to produce an optically clear solution.

Size-selective Precipitation. Nanocrystallites were dispersed in a solution of ~10% butanol in hexane. Methanol was then added dropwise to this stirring solution until opalescence persisted. Separation of supernatant and flocculate by centrifugation produced a precipitate enriched with the largest crystallites in the sample. This procedure was repeated until no further sharpening of the optical absorption spectrum was noted. Size-selective precipitation can be carried out in a variety of solvent/nonsolvent pairs, including pyridine/hexane and chloroform/methanol.

Surface Exchange. Crystallite surface derivatization can be modified by repeated exposure to an excess of a competing capping group. Heating to ~60 °C a mixture of ~50 mg of TOPO/TOP capped crystallites and 5-10 mL of pyridine gradually dispersed the crystallites in the solvent. Treatment of the dispersion with excess hexane resulted in the flocculation of the crystallites which are then isolated by centrifugation. The process of dispersion in pyridine and flocculation with hexane was repeated a number of times to produce crystallites which dispersed readily in pyridine, methanol, and aromatics but no longer dispersed in aliphatics.

Example 2. Preparation of CdSe. A second route to the production of CdSe core replaces the phosphine chalcogenide precursors in Example 1 with (TMS)₂Se. The smallest (~12 Å) CdSe species are produced under milder conditions with injection and growth carried out at ~100 °C. The product was further treated as described in Example 1.

Example 3. Preparation of (CdSe)/ZnS. Nearly monodisperse CdSe quantum dots ranging from 23 Å to 55 Å in diameter were synthesized and purified via size-selective precipitation as described in Example 1.

5 A flask containing 5g of TOPO was heated to 190 °C under vacuum for several hours then cooled to 60 °C after which 0.5 mL trioctylphosphine (TOP) was added. Roughly 0.1-0.4 μmols of CdSe dots dispersed in hexane were transferred into the reaction vessel via syringe and the solvent was pumped off.

Diethyl zinc (ZnEt_2) and hexamethyldisilathiane ($(\text{TMS})_2\text{S}$) were used as the Zn and S precursors, respectively. The amounts of Zn and S precursors needed to grow a
10 ZnS shell of desired thickness for each CdSe sample were determined as follows: First, the average radius of the CdSe dots was estimated from TEM or SAXS measurements. Next, the ratio of ZnS to CdSe necessary to form a shell of desired thickness was calculated based on the ratio of the shell volume to that of the core assuming a spherical core and shell and taking into account the bulk lattice parameters of CdSe and ZnS. For
15 larger particles the ratio of Zn to Cd necessary to achieve the same thickness shell is less than for the smaller dots. The actual amount of ZnS that grows onto the CdSe cores was generally less than the amount added due to incomplete reaction of the precursors and to loss of some material on the walls of the flask during the addition.

Equimolar amounts of the precursors were dissolved in 2-4 mL TOP inside an
20 inert atmosphere glove box. The precursor solution was loaded into a syringe and transferred to an addition funnel attached to the reaction flask. The reaction flask containing CdSe dots dispersed in TOPO and TOP was heated under an atmosphere of N_2 . The temperature at which the precursors were added ranged from 140 °C for 23 Å diameter dots to 220 °C for 55 Å diameter dots. When the desired temperature was
25 reached the Zn and S precursors were added dropwise to the vigorously stirring reaction mixture over a period of 5-10 minutes.

After the addition was complete the mixture was cooled to 90 °C and left stirring for several hours. Butanol (5mL) was added to the mixture to prevent the TOPO from solidifying upon cooling to room temperature. The overcoated particles were stored in
30 their growth solution to ensure that the surface of the dots remained passivated with

TOPO. They were later recovered in powder form by precipitating with methanol and redispersing into a variety of solvents including hexane, chloroform, toluene, THF and pyridine.

5 In some cases, the as-grown CdSe crystallites were judged to be sufficiently monodisperse that no size-selective precipitation was performed. Once these CdSe particles had grown to the desired size, the temperature of the reaction flask was lowered and the Zn and S precursors were added dropwise to form the overcapping.

Optical Characterization. UV-Visible absorption spectra were acquired on an HP 8452 diode array spectrophotometer. Dilute solutions of dots in hexane were placed in 1
10 cm quartz cuvettes and their absorption and corresponding fluorescence were measured. The photoluminescence spectra were taken on a SPEX Fluorolog-2 spectrometer in front face collection mode. The room temperature quantum yields were determined by comparing the integrated emission of the dots in solution to the emission of a solution of rhodamine 590 or rhodamine 640 of identical optical density at the excitation
15 wavelength.

What is claimed is:

1 1. A coated nanocrystal capable of light emission, comprising:
2 a core selected from the group consisting of CdX, where x = S, Se, Te, and
3 mixtures thereof, said core being a member of a substantially monodisperse particle
4 population; and
5 an overcoating of ZnY, where Y = S, Se, uniformly deposited thereon, said coated
6 core characterized in that when irradiated the particles emit light in a narrow spectral
7 range of no greater than about 60 nm at full width half max (FWHM).

8
9 2. A coated nanocrystal capable of light emission, comprising:
10 a core selected from the group consisting of CdX, where x = S, Se, Te, and
11 mixtures thereof, said core being a member of a substantially monodisperse particle
12 population; and
13 an overcoating of ZnY, where Y = S, Se, uniformly deposited thereon, said coated
14 core characterized in that the nanocrystal exhibits less than a 10% rms deviation in
15 diameter of the core.

16
17 3. The coated nanocrystal of claim 1, wherein the spectral range is not
18 greater than about 40 nm at full width half max (FWHM).

19
20 4. The coated nanocrystal of claim 1, wherein the spectral range is not
21 greater than about 30 nm at full width half max (FWHM).

22
23 5. The coated nanocrystal of claim 1, wherein the coated nanocrystal exhibits
24 photoluminescence having quantum yields of greater than 30%.

25
26 6. The coated nanocrystal of claim 1, wherein the coated nanocrystal exhibits
27 photoluminescence having quantum yields in the range of about 30 to 50%.

28

1 7. The coated nanocrystal of claim 2, wherein the coated nanocrystal
2 exhibits less than a 5% rms deviation in size of the core.

3
4 8. The coated nanocrystal of claim 1 or 2, wherein the overcoating comprises
5 one to two monolayers of ZnY.

6
7 9. The coated nanocrystal of claim 1, wherein the narrow spectral range is
8 selected from the spectrum in the range of about 470 nm to about 620 nm.

9
10 10. The coated nanocrystal of claim 2, wherein the particle size of the core is
11 selected from the range of about 20Å to about 125 Å.

12
13 11. The coated nanocrystal of claim 1 or 2, wherein the nanocrystal further
14 comprises an organic layer on the nanocrystal outer surface.

15
16 12. The coated nanocrystal of claim 11, wherein the organic layer is
17 comprised of moieties selected to provide compatibility with a suspension medium.

18
19 13. The coated nanocrystal of claim 11, wherein the organic layer is
20 comprised of moieties selected to exhibit affinity for the outer surface of the
21 nanocrystal.

22
23 14. The coated nanocrystal of claim 13, wherein the organic layer comprises
24 a short-chain polymer terminating in a moiety having affinity for a suspending
25 medium.

26
27 15. A method of preparing a coated nanocrystal capable of light emission,
28 comprising:

1 introducing a substantially monodisperse first semiconductor core population and
2 a precursor capable of thermal conversion into a second semiconductor material into a
3 coordinating solvent,

4 wherein the coordinating solvent is maintained at a temperature sufficient to
5 convert the precursor into the second semiconductor material yet insufficient to
6 substantially alter the monodispersity of the first semiconducting core,

7 wherein the second semiconductor material has a band gap greater than the first
8 semiconducting nanocrystal, and

9 whereby an overcoating of the second semiconductor material is formed on the
10 first semiconducting nanocrystal.

11
12 16. The method of claim 15, further comprising:
13 monitoring the monodispersity of the nanocrystal during conversion of the
14 precursor and overcoating of the first semiconductor nanocrystal.

15
16 17. The method of claim 15, wherein the is was lowered in response to a
17 spreading of the size distribution as estimated from the absorption spectra.

18
19 18. The method of claim 15, wherein the temperature is increased in response
20 to when monitoring indicates growth appears to stop.

21
22 19. The method of claim 15, wherein the first semiconductor crystal is
23 selected from the group consisting of CdX, where X = S, Se and Te.

24
25 20. The method of claim 15, wherein the second semiconductor material is
26 selected from the group consisting of ZnS, ZnSe, CdS and CdSe and mixtures thereof.

27
28 21. The method of claim 15, wherein the particle size of the core is in the
29 range of about 20Å to about 125 Å.

1 22. The method of claim 15, wherein the nanocrystal further comprises an
2 organic layer on the nanocrystal outer surface.

3

4 23. The method of claim 22, wherein the organic layer is obtained by
5 exposing the nanocrystal to an organic compound having affinity for the nanocrystal
6 surface, whereby the organic compound displaces the coordinating solvent.

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HIGHLY LUMINESCENT COLOR-SELECTIVE MATERIALS

DS1.349876.1

Abstract of the Invention

A coated nanocrystal capable of light emission includes a substantially monodisperse nanoparticle selected from the group consisting of CdX, where x = S, Se, Te and an overcoating of ZnY, where Y = S, Se, uniformly deposited thereon, said coated nanoparticle characterized in that when irradiated the particles exhibit photoluminescence in a narrow spectral range of no greater than about 60 nm, and most preferably 40 nm, at full width half max (FWHM). The particle size of the nanocrystallite core is in the range of about 20Å to about 125 Å, with a deviation of less than 10% in the core. The coated nanocrystal exhibits photoluminescence having quantum yields of greater than 30%.

DS1.349876.2

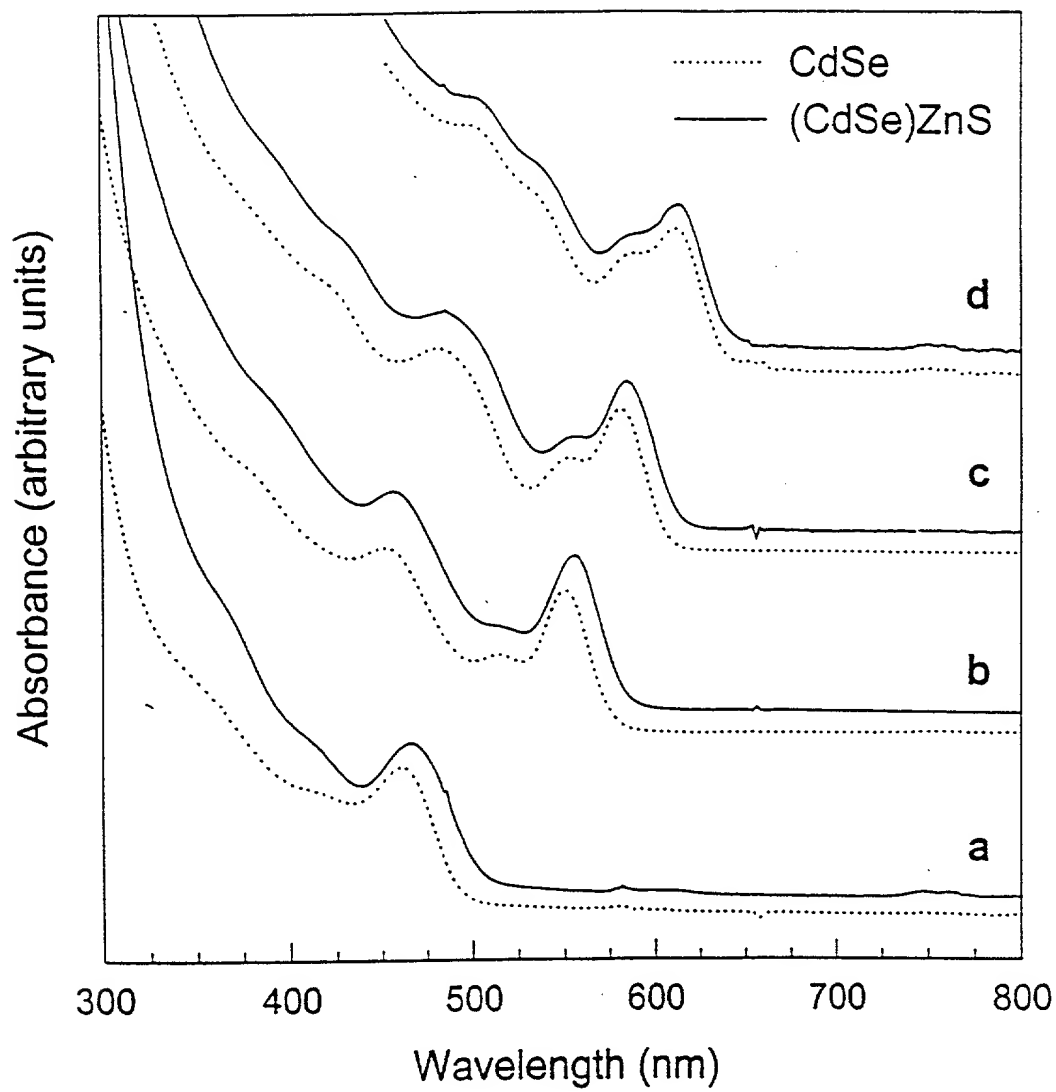


Figure 1

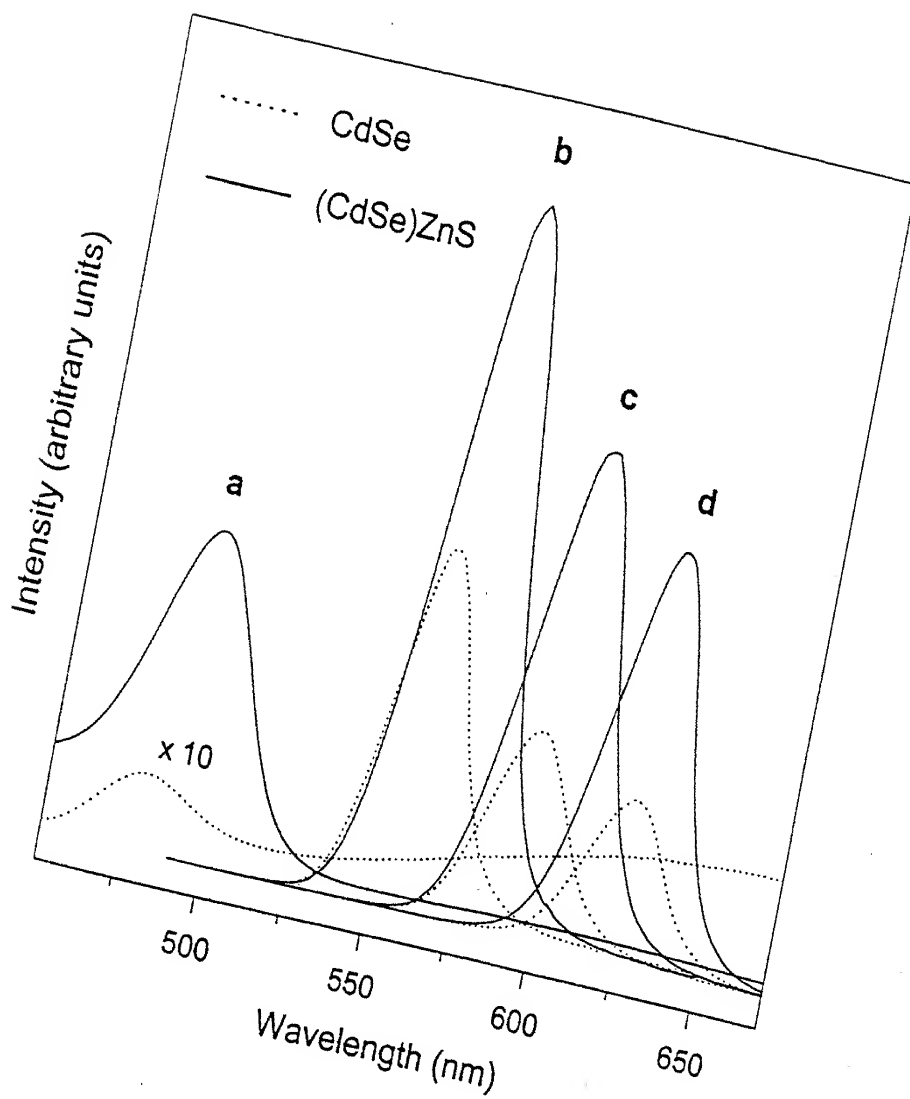


Figure 2

Felice Frankel

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Figure 3

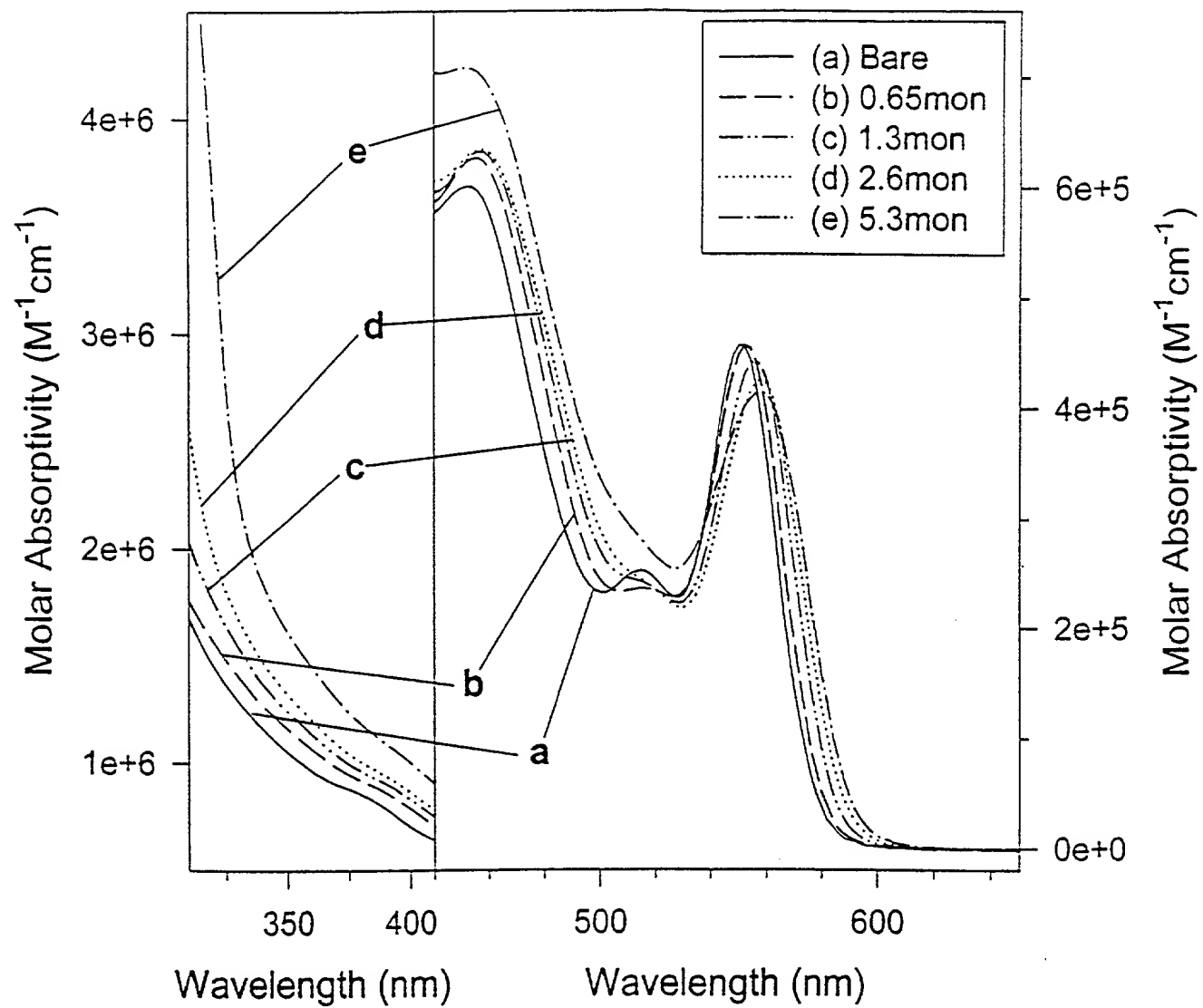


Figure 4

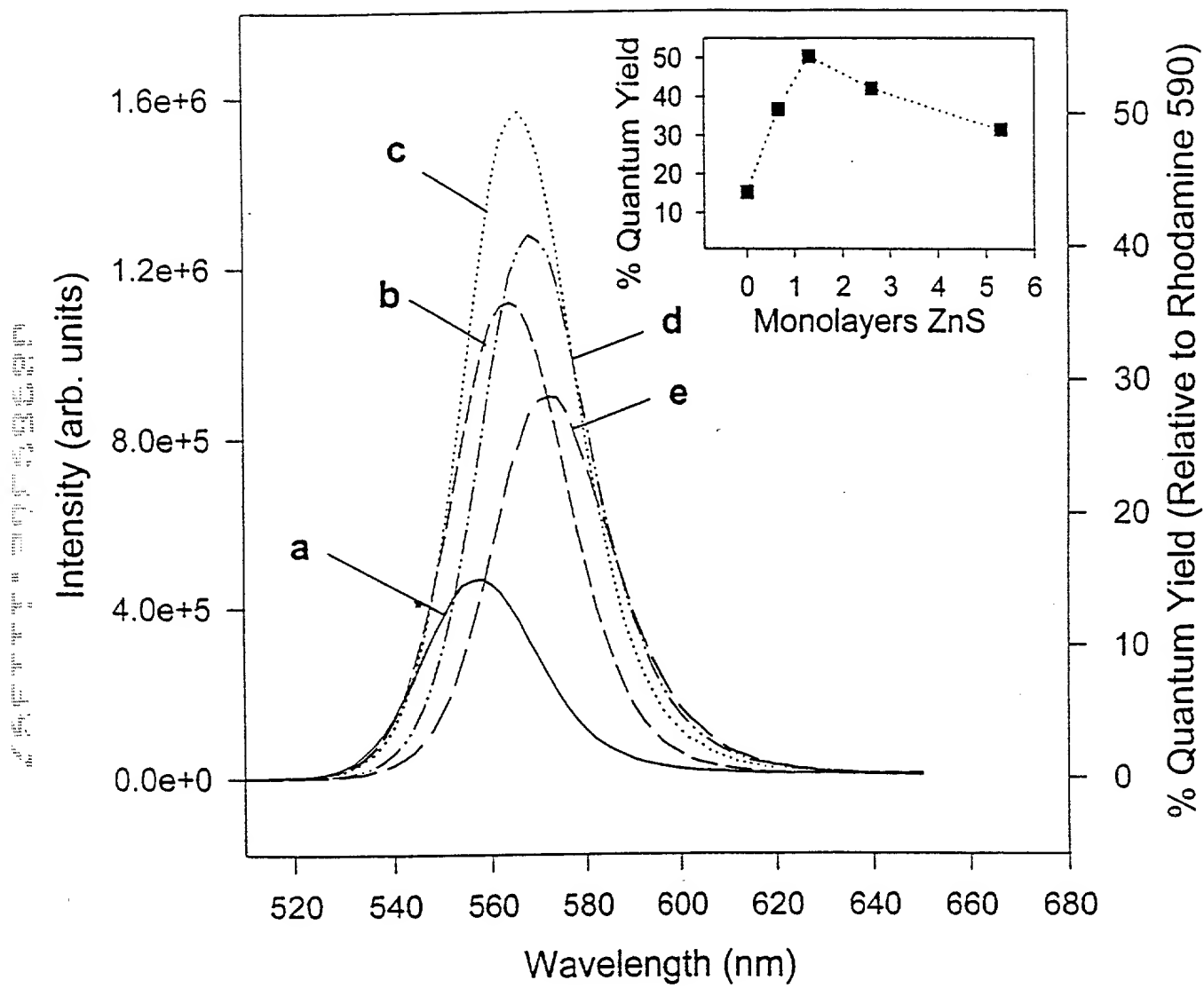


Figure 5

Applicant: Bawendi et al. Examiner:
Serial No.: Art Unit:
Filed: November 13, 1997 Attorney Docket No.: 0492611-0289
For: HIGHLY LUMINESCENT COLOR-SELECTIVE MATERIALS

VERIFIED STATEMENT (DECLARATION) CLAIMING SMALL ENTITY
STATUS (37 CFR 1.9 (F) AND 1.27 (D)) - SMALL BUSINESS CONCERN

I hereby declare that I am:

- () the owner of the small business concern identified below:
(x) an official of the small business concern empowered to act on behalf of the concern identified below:

NAME OF CONCERN: Massachusetts Institute of Technology
ADDRESS OF CONCERN: 77 Massachusetts Avenue, Cambridge, MA 02139

I hereby declare that the above-identified small business concern qualifies as a small business concern as defined in 13 CFR 121.3-18, and reproduced in 37 CFR 1.9(d), for purposes of paying reduced fees under section 41(a) and (b) of Title 35, United States Code, in that the number of employees of the concern, including those of its affiliates, does not exceed 500 persons. For purposes of this statement, (1) the number of employees of the business concern is the average over the previous fiscal year of the concern of the persons employed on a full-time, part-time or temporary basis during each of the pay periods of the fiscal year, and (2) concerns are affiliates of each other when either, directly or indirectly, one concern controls or has the power to control the other, or a third party or parties controls or has the power to control both.

I hereby declare that rights under contract or law have been conveyed to and remain with the small business concern identified above with regard to the invention, entitled HIGHLY LUMINESCENT COLOR-SELECTIVE MATERIALS by Mounji Bawendi, Klavs Jensen, Bashire Dabbousi, Xavier Rodriguez-Viejo and Frederic Victor Mikulec described in

- (XX) the specification filed herewith
() application serial no. 08/808,120, filed February 28, 1997
() patent no. , issued

If the rights of the above-identified small business concern are not exclusive, each individual, concern or organization having rights to the invention is listed below* and no rights to the invention are held by any person, other than the inventor(s), who could not qualify as a small business concern under 37 CFR 1.9(d) or by any concern which would not qualify as a small business concern under 37 CFR 1.9(d) or a nonprofit organization under 37 CFR 1.9(e). *NOTE: Separate verified statements are required from each named

person, concern or organization having rights to the invention averring to their status as small entities. (37 CFR 1.27)

FULL NAME _____

ADDRESS _____

() Individual () Small Business Concern () Nonprofit Organization

FULL NAME _____

ADDRESS _____

() Individual () Small Business Concern () Nonprofit Organization

I acknowledge the duty to file, in this application or patent, notification of any change in status resulting in loss of entitlement to small entity status prior to paying, or at the time of paying, the earliest of the issue fee or any maintenance fee due after the date on which status as a small entity is no longer appropriate. (37 CFR 1.28(b))

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application, any patent issuing thereon, or any patent to which this verified statement is directed.

NAME OF PERSON SIGNING:

TITLE OF PERSON:

ADDRESS OF PERSON: 77 Massachusetts Avenue, Cambridge, MA 02139

SIGNATURE _____ DATE _____

**DECLARATION AND POWER OF ATTORNEY
FOR PATENT APPLICATION**

As below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name. I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled HIGHLY LUMINESCENT COLOR-SELECTIVE MATERIALS, the specification of which was filed on November 13, 1997, as United States Serial No.

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, §1.56.

I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith: Mary Rose Scozzafava, Reg. No. 36,268; Sam Pasternack, Reg. No. 29,576; Mary Raynor Jimenez, Reg. No. 37,219; David J. Powsner, Reg. No. 31,868; Brenda H. Jarrell, Reg. No. 39,223; Douglas D. Robinow, Reg. No. 40,787; and Kevin M. Tormey, Reg. No. 41,351.

Address all telephone calls to Mary Rose Scozzafava at (617) 248-5138.

Address all correspondence to Mary Rose Scozzafava at Choate, Hall & Stewart, Exchange Place, 53 State Street, Boston, Massachusetts 02109.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of sole or first inventor: Moungi Bawendi

Inventor's signature _____ Date _____

Residence: 4 Trowbridge P. #3F, Cambridge, Massachusetts 02178

Citizenship: United States of America

Post Office Address: same

Full name of second inventor: Klavs F. Jensen
Inventor's signature _____ Date _____
Residence: 103 Outlook Drive, Lexington, Massachusetts 02173
Citizenship: United States of America
Post Office Address: same

Full name of third inventor: Bashire O. Dabbousi
Inventor's signature _____ Date _____
Residence: _____
Citizenship: _____
Post Office Address: _____

Full name of fourth inventor: Xavier Rodriguez-Viejo
Inventor's signature _____ Date _____
Residence: Cami de Can Gatzet 57 3-1, 08190 Sant Cugat del Valles, Spain
Citizenship: Spain
Post Office Address: same

Full name of fifth inventor: Frederic Victor Mikulec
Inventor's signature _____ Date _____
Residence: 96 Willow Avenue, Apt. 2, Somerville, Massachusetts 02144
Citizenship: United States of America
Post Office Address: same